



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service



APR -6 1998

Food and Drug Administration
Rockville MD 20857

Re: GLYSET®
Docket No. 97E-0077
#46

Stephen G. Kunin
Deputy Assistant Commissioner for
Patent Policy and Projects
Office of the Assistant Commissioner for Patents
U.S. Patent and Trademark Office
Crystal Park Building 2, Suite 919
Washington, DC 20231

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PATENT EXTENSION
A/C PATENTS

Dear Mr. Kunin:

This is in regard to the patent term extension application for U.S. Patent No. 4,639,436 filed by Bayer Aktiengesellschaft under 35 U.S.C. § 156. The patent claims the human drug product GLYSET® (miglitol), New Drug Application NDA 20-682.

In the July 18, 1997, issue of the Federal Register (62 Fed. Reg. 38561), the Food and Drug Administration published its determination of this product's regulatory review period, as required under 35 U.S.C. § 156(d)(2)(A). The notice provided that on or before January 14, 1998, 180 days after the publication of the determination, any interested person could file a petition with FDA under 35 U.S.C. § 156(d)(2)(B)(i) for a determination of whether the patent term extension applicant acted with due diligence during the regulatory review period.

The 180-day period for filing a due diligence petition pursuant to this notice has expired and FDA has received no such petition. Therefore, FDA considers the regulatory review period determination to be final.

Please let me know if we can provide further assistance.

Sincerely yours,

Ronald L. Wilson, Director
Health Assessment Policy Staff
Office of Health Affairs

cc: Jeffrey M. Greenman, Esq.
Bayer Corporation
400 Morgan Lane
West Haven, CT 06516